Supplemental Reply to Office Action of March 11, 2008

AMENDMENTS TO THE CLAIMS

1-15. (Canceled)

16. (Currently Amended) A DNA synthesis reaction composition comprising:

1) a DNA polymerase;

2) water-soluble acidic macromolecular substances or water-soluble salts thereof,

wherein said water-soluble acidic macromolecular substances are one or more substances

selected from the group consisting of sulfated-fucose-containing polysaccharides polysaccharide

F, sulfated-fucose-containing polysaccharide-U, hyaluronic acid, and alginic acid, polyglutamic

acids, polyacrylic acids and polystyrene sulfates; and

3) components necessary for DNA synthesis using DNA polymerase.

17. (Canceled)

18. (Currently Amended) A DNA synthesis reaction composition comprising:

1) two or more kinds of DNA polymerases;

2) water-soluble acidic macromolecular substances or water-soluble salts thereof,

wherein said water-soluble acidic macromolecular substances are one or more substances

selected from the group consisting of sulfated-fucose-containing polysaccharides polysaccharide-

F. sulfated-fucose-containing polysaccharide-U, dermatan sulfate (chondroitin sulfate B),

hyaluronic acid, and alginic acid, polyglutamic acids, polyacrylic acids, polyvinyl sulfates and

polystyrene sulfates; and

Supplemental Reply to Office Action of March 11, 2008

3) components necessary for DNA synthesis using DNA polymerase,

wherein the two or more kinds of DNA polymerases comprise a DNA polymerase having

3'→5' exonuclease activity, and a DNA polymerase having no 3'→5' exonuclease activity.

19-30. (Canceled)

31. (Currently Amended) A kit for use in in vitro DNA synthesis, wherein the kit

comprises:

1) a DNA polymerase;

2) a reaction buffer comprising water-soluble acidic macromolecular substances or water-

soluble salts thereof, wherein said water-soluble acidic macromolecular substances are one or

more substances selected from the group consisting of sulfated-fucose-containing

polysaccharides polysaccharide-F, sulfated-fucose-containing polysaccharide-U, hyaluronic acid,

and alginic acid, polyglutamic acids, polyacrylic acids and polystyrene sulfates; and

3) dNTP, wherein N is a mixture of adenine, thymine, guanine and cytosine.

32-33. (Canceled)

34. (Previously Presented) The kit according to claim 31, wherein said DNA polymerase

is a thermostable DNA polymerase.

35. (Canceled)

Supplemental Reply to Office Action of March 11, 2008

36. (Currently Amended) A kit for use in in vitro DNA synthesis, wherein the kit

comprises:

1) two or more kinds of DNA polymerases, wherein the two or more kinds of DNA

polymerases comprise a DNA polymerase having 3'→5' exonuclease activity, and a DNA

polymerase having no 3'→5' exonuclease activity

2) a reaction buffer comprising water-soluble acidic macromolecular substances or water-

soluble salts thereof, wherein said water-soluble acidic macromolecular substances are one or

more substances selected from the group consisting of sulfated-fucose-containing

polysaccharides polysaccharide-F, sulfated-fucose-containing polysaccharide-U, dermatan

sulfate (chondroitin sulfate B), hyaluronic acid, and alginic acid, polyglutamic acids, polyacrylic

acids, polyvinyl sulfates and polystyrene sulfates; and

3) dNTP, wherein N is a mixture of adenine, thymine, guanine and cytosine.

37. (Canceled)

38. (Previously Presented) The kit according to claim 36, wherein at least one of said two

or more kinds of DNA polymerases is thermostable.

39. (Canceled)

Supplemental Reply to Office Action of March 11, 2008

40. (Previously Presented) The DNA synthesis reaction composition according to

claim 16, wherein said water-soluble acid macromolecular substances or water-soluble salts

thereof are present in the composition at about 0.1 ng to about 5 ug, and wherein the composition

is about 50 µl in total volume.

41. (Previously Presented) The DNA synthesis reaction composition according to

claim 16, wherein said DNA polymerase is selected from the group consisting of: pol I-type

DNA polymerase, E. coli DNA polymerase I, Klenow fragment, Thermococcus aquaticus-

derived DNA polymerase, α-type DNA polymerase, α-type Pyrococcus furiosus-derived DNA

polymerase, Thermococcus litralis-derived DNA polymerase and Pyrococcus sp.-derived DNA

polymerase.

42. (Previously Presented) The DNA synthesis reaction composition according to

claim 16, wherein said DNA polymerase is selected from the group consisting of: E. coli DNA

polymerase I, Klenow fragment, Taq DNA polymerase, VENT DNA polymerase, Pyrobest DNA

polymerase, Pfu DNA polymerase I, Pfu DNA polymerase II, Ex-Taq DNA polymerase, KOD

dash DNA polymerase, DEEP VENT DNA polymerase, KOD DNA polymerase and LA-Taq

DNA polymerase.

43-44. (Canceled)

Supplemental Reply to Office Action of March 11, 2008

45. (Previously Presented) The DNA synthesis reaction composition according to claim

18, wherein said two or more kinds of DNA polymerases are selected from the group consisting

of: pol I-type DNA polymerase, E. coli DNA polymerase I, Klenow fragment, Thermococcus

aquaticus-derived DNA polymerase, α-type DNA polymerase, α-type Pyrococcus furiosus-

derived DNA polymerase. Thermococcus litralis-derived DNA polymerase and Pyrococcus sp.-

derived DNA polymerase.

46. (Previously Presented) A method of enhancing DNA synthesis which comprises:

incubating the synthesis reaction composition of claim 16 in the presence of a nucleic

acid to be amplified.

47. (Previously Presented) A method of enhancing DNA synthesis which comprises:

incubating the synthesis reaction composition of claim 18 in the presence of a nucleic

acid to be amplified.